IN THE CLAIMS

Please amend the claims as follows:

Claim 1 (Currently Amended): A method of treating a human patient suffering from an ocular allergy, comprising:

administering to said patient an ophthalmic composition eontaining comprising from about 0.01% to about 0.1% of a macrolide compound.

Claim 2 (Currently Amended): A method according to claim 1, wherein said ocular allergy is allergic conjunctivitis.

Claim 3 (Currently Amended): A method according to claim 1 or 2, wherein said composition comprises from about 0.03% to about 0.06% of said macrolide compound.

Claim 4 (Currently Amended): A method according to claim 3, wherein said macrolide compound ophthalmic composition contains comprises about 0.03% of said macrolide compound.

Claim 5 (Currently Amended): A method according to claim 1, wherein said macrolide compound is FK506.

Claim 6 (Currently Amended): A method according to claim 1, wherein said ophthalmic composition is <u>an</u> eye drop.

Claim 7 (Currently Amended): A method according to claim 6, wherein said eye drop further contains comprises polyvinyl alcohol.

Claim 8 (Currently Amended): A method according to claim 7, wherein said eye drop eontains comprises about 0.03% of said macrolide compound.

Claim 9 (Original): A method according to claim 8, wherein said eye drop is administered from about one to about 4 times per day.

Claim 10 (Currently Amended): A method according to any of claims 1 to 9 claim 1, wherein said macrolide compound is a compound having the following formula (I) or a pharmaceutically acceptable salt thereof:

$$R^{24}$$
 R^{6}
 R^{19}
 R^{10}
 R^{23}
 R^{10}
 R^{10}

wherein adjacent pairs of R¹ and R², R³ and R⁴, and R⁵ and R⁶ each independently a) consist of two adjacent hydrogen atoms, wherein R² is optionally alkyl, or

b) form another bond optionally between carbon atoms binding with the members of said pairs;

 R^7 is hydrogen atom, hydroxy, alkyloxy or protected hydroxy, or may form oxo with R^1 ;

R⁸ and R⁹ each independently show hydrogen atom or hydroxy;

R¹⁰ is hydrogen atom, alkyl, alkyl substituted by one or more hydroxy, alkenyl, alkenyl substituted by one or more hydroxy or alkyl substituted by oxo;

X is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula -CH₂O-;

Y is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula N-NR¹¹R¹² or N-OR¹³;

R¹¹ and R¹² each independently show hydrogen atom, alkyl, aryl or tosyl;

 R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{22} and R^{23} each independently show hydrogen atom or alkyl;

 R^{24} is an optionally substituted ring that may contain one or more hetero atom(s); and n is 1 or 2.

Claim 11 (Original): A method according to claim 10, wherein said macrolide compound has the following structure:

Claim 12 (Currently Amended): An ophthalmic composition for treatment of ocular allergy eontaining comprising from about 0.01% to about 0.1% of a macrolide compound.

Claim 13 (Currently Amended): An ophthalmic composition according to claim 12, wherein said ocular allergy is allergic conjunctivitis.

Claim 14 (Currently Amended): An ophthalmic composition according to claim 12 or 13, comprising which contains from about 0.03% to about 0.06% of said macrolide compound.

Claim 15 (Currently Amended): An ophthalmic composition according to claim 14, comprising which contains about 0.03% of said macrolide compound.

Claim 16 (Currently Amended): An ophthalmic composition according to claim 12, wherein said macrolide compound is FK506.

Claim 17 (Currently Amended): An ophthalmic composition according to claim 12, which is an eye drop.

Claim 18 (Currently Amended): An ophthalmic composition according to claim 17, wherein said eye drop further contains comprises polyvinyl alcohol.

Claim 19 (Currently Amended): An ophthalmic composition according to claim 18, wherein said eye drop contains comprises about 0.03% of said macrolide compound.

Claim 20 (Original): An ophthalmic composition according to claim 19, wherein said eye drop is administered from about one to about 4 times per day.

Claim 21 (Currently Amended): An ophthalmic composition according to any of claims 12 to 20 claim 12, wherein said macrolide compound is a compound having the following formula (I) or a pharmaceutically acceptable salt thereof:

wherein adjacent pairs of R¹ and R², R³ and R⁴, and R⁵ and R⁶ each independently a) consist of two adjacent hydrogen atoms, wherein R² is optionally alkyl, or

b) form another bond optionally between carbon atoms binding with the members of said pairs;

 R^7 is hydrogen atom, hydroxy, alkyloxy or protected hydroxy, or may form oxo with R^1 ;

R⁸ and R⁹ each independently show hydrogen atom or hydroxy;

R¹⁰ is hydrogen atom, alkyl, alkyl substituted by one or more hydroxy, alkenyl, alkenyl substituted by one or more hydroxy or alkyl substituted by oxo;

X is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula -CH₂O-;

Y is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula N-NR¹¹R¹² or N-OR¹³;

R¹¹ and R¹² each independently show hydrogen atom, alkyl, aryl or tosyl;

 R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{22} and R^{23} each independently show hydrogen atom or alkyl;

 R^{24} is an optionally substituted ring that may contain one or more hetero atom(s); and n is 1 or 2.

Claims 22-33 (Canceled).

Claim 34 (Currently Amended): A commercial package comprising the ophthalmic composition of any of claims 12 to 22 claim 12 and a written matter associated therewith, the written matter stating that the composition can or should be used for allergic conjunctivitis.